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(54) **Topical azithromycin compositions for the treatment of ocular infections**

(57) The invention relates to topical compositions

containing azithromycin for the treatment of ocular infections.

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DescriptionField of the invention

5 [0001] This invention relates to methods of treating eye infections by topically administering azithromycin to an eye of an animal in need of such treatment.

Background of the invention

10 [0002] Azithromycin is the U.S.A.N. (generic name) for 9a-aza-9a-methyl-9-deoxo-9a-homoerythromycin A, a broad spectrum antimicrobial compound derived from erythromycin A. Azithromycin is disclosed in US patent no. 4,474,768 to Bright and US patent no. 4,517,359 to Kobrehel et al. These patents disclose that azithromycin and certain derivatives thereof possess antibacterial properties and are accordingly useful as antibiotics.

15 [0003] Azithromycin is commonly administered orally, in a number of different dosage forms such as tablets, capsules, and suspensions, for the treatment of antibacterial infections. Until the present invention, however, azithromycin was not known to be effective when topically administered to the eye. Azithromycin is known to be effective for the treatment of eye infections in humans when administered systemically, e.g., orally. However, it is also known that antibiotics which are effective when administered by a systemic route are not necessarily effective when applied topically, directly to the eye. For example, it has been reported that when tetracyclines are applied to the cornea, they do not penetrate the
20 intact normal cornea even though they are able to diffuse into spinal fluid and into ocular fluids if the systemic dose is high enough (Douvas MG, et al, Arch Ophthalmol. 46:57, 1951).

Summary of the invention

25 [0004] This invention provides a method of treating an ocular infection, comprising topically administering, to an eye of an animal, including man, in need of such treatment, an ocular infection-treating amount of azithromycin. Topical administration means the application, directly to the surface of an eye, of azithromycin in a composition comprising azithromycin and a pharmaceutically acceptable topical carrier. In a preferred embodiment, the composition is applied directly to an eye as a single dose (equivalent to approximately 5mg) per day for five days. It is noted that a "single
30 dose" means the amount for a single eye.

[0005] The invention further provides pharmaceutical compositions for topical application directly to an eye of an animal, including a human, said compositions being suitable for the treatment of an ocular infection, comprising azithromycin and a pharmaceutical vehicle suitable for topical application, wherein said azithromycin is at a concentration in said vehicle sufficient to remediate said ocular infection. In a preferred embodiment, the concentration of azithromycin
35 in the vehicle is such that a single dose (approx. 5mg per eye) of said composition administered once daily for five days remedies the infection. The capability of achieving once-a-day topical dosing with azithromycin was highly unexpected considering that most drugs are rapidly cleared from the precorneal area by tear drainage. Thus, commonly most topical regimens using known antibiotics such as gentamycin and erythromycin must be administered frequently with application rates of 4-6 times daily sometimes being required to produce effective drug levels in target ocular
40 tissues. Topical formulations of azithromycin, by contrast, achieve relatively high and sustained levels in ocular tissues including cornea, conjunctiva, lids, and sclera. Because of the excellent penetration of azithromycin into ocular tissues, patient compliance is expected to be significantly enhanced by virtue of this invention.

[0006] The types of ocular infections treatable through the topical administration of azithromycin broadly include any eye infection caused by a bacterial species known to be amenable to systemic treatment with azithromycin. In particular,
45 the invention is applicable to the treatment of trachoma, a chronic follicular conjunctivitis due to *Chlamydia trachomatis*, the world's leading cause of preventable blindness.

Detailed description

50 [0007] The term "azithromycin" includes the pharmaceutically acceptable salts thereof, and anhydrous as well as hydrated forms. The azithromycin is preferably present as the dihydrate, disclosed, for example, in published European Patent Application 0 298 650 A2 and in co-pending U. S. application 07/994,040 filed December 21, 1992, each of which is herein incorporated by reference.

[0008] Compositions (sometimes referred to herein as "formulations") according to the invention comprise azithromycin and a pharmaceutically acceptable vehicle suitable for topical application to an eye. Azithromycin (calculated using the dihydrate) is typically present in the composition at a concentration of 0.1 to 2.5 weight % (w/w), usually 0.2 to 2.0 weight %, based on the weight of the composition. A preferred concentration is 0.5 weight %.

[0009] The compositions can include a preservative if desired, although preferred compositions do not contain a

preservative. The compositions can also optionally contain surfactants, viscosity enhancers, buffers, sodium chloride, and water to form aqueous sterile ophthalmic solutions and suspensions. In order to prepare sterile ophthalmic ointment formulations, azithromycin is combined with an appropriate vehicle such as mineral oil, liquid lanolin, or white petrolatum. Sterile ophthalmic gel formulations containing azithromycin can be prepared by suspending azithromycin in a hydrophilic base prepared from a combination of, for example, a carboxyvinyl polymer sold under the designation Carbopol® (registered trademark of the B. F. Goodrich Company for a series of such polymers) according to published formulations for analogous ophthalmic preparations. Tonicity agents may also be incorporated in such gel formulations.

[0010] Azithromycin can be formulated as an ophthalmic solution in isotonic saline using glycerine as an isotonicity agent. A preservative can optionally be included as an excipient. Such ophthalmic solutions also include a pharmaceutically acceptable buffering agent, typically a combination of boric acid and sodium borate, sufficient to maintain the pH of the solution between 7 and 8.

[0011] A preferred composition is 0.5 % w/w azithromycin dihydrate suspended in an inert, non-allergenic, preservative-free vehicle consisting of white petrolatum (55% w/w), mineral oil (42.5% w/w), and lanolin (2% w/w).

[0012] The invention is further disclosed by means of the following non-limiting examples. In the examples, reference to "water" means sterile water, suitable for use as water for injection.

Example 1

[0013] A preferred embodiment was prepared by incorporating 5g of azithromycin dihydrate into 995 g of a sterile vehicle comprising 55% by weight of petrolatum, 43% by weight of light mineral oil, and 2% by weight of lanolin. The procedure involved first heating an excess amount of the ointment vehicle of the aforementioned composition to 70°C in a glass vessel to produce a melt. In the next step, 995g of the molten sterile ointment was transferred into a compounding vessel equipped with a mixer and 5g of the azithromycin dihydrate was added to the melt under agitation at 70°C to form a suspension. The azithromycin containing ointment was rapidly cooled by placing the compounding vessel in an ice bath. The "0.5% azithromycin ointment" was then filled into 1cc unit dose plastic syringes for dose application. This ointment may be sterilized by gamma radiation using a cobalt-60 source.

Example 2

[0014] A 0.5 weight percent azithromycin dihydrate ophthalmic solution is prepared by dissolving 50g of azithromycin dihydrate (0.5 weight %), 67.0 g (0.67 weight percent) boric acid, 20.7 g (0.207 weight percent) sodium borate decahydrate, 100 g (1.0 weight percent) glycerin, 100 g of polyethylene glycol 300 (1.0 weight percent), and 0.40 g (0.004 weight percent) thimerosal (as a preservative) in about 8000 g of deionized distilled water. The pH is adjusted to 7.2 with HCl and NaOH. The final batch weight is brought to 10,000 g with the addition of the required amount of water. The final solution is filtered through a 0.2 micron Millipore filter and filtered into vials.

Example 3

[0015] In a preferred embodiment, an approximate 0.5 weight percent azithromycin dihydrate ophthalmic suspension is prepared as follows: 600 g of petrolatum is heated to 90°C for 2 hours in a jacketed 316 stainless steel vessel. The temperature is then decreased to 60°C. Light mineral oil, 350 g, is added to the petrolatum under mild agitation. The solution is passed through a sintered glass filter. Azithromycin dihydrate, 5g, is dispersed into the mineral oil/petrolatum solution under agitation to form a finely dispersed suspension. The suspension is cooled under slow agitation to form a semisolid suspension. The suspension is filled into plastic, polypropylene tubes and sterilized by gamma radiation using a cobalt-60 source.

Example 4

[0016] A 0.5 weight percent azithromycin dihydrate ophthalmic suspension is prepared as follows: 600 g of PEG 4000 is heated to 90°C for 2 hours in a jacketed 316 stainless steel vessel. The temperature is brought down to 60°C. PEG 400, 350 g, is added to the petrolatum under mild agitation. The solution is passed through a sintered glass filter. Azithromycin dihydrate, 50 g, is dispersed into the PEG 4000/PEG 400 solution under agitation to form a finely dispersed suspension. The suspension is cooled under slow agitation to form a semisolid suspension. The suspension is filled into plastic, polypropylene tubes and sterilized by gamma radiation using a cobalt-60 source.

EXAMPLE 5

[0017] The following gel is prepared under strictly aseptic conditions:

	% w/w
Azithromycin Dihydrate	3.50
Chlorbutol BP	0.50
Carbopol® 934P	2.50
NaOH (4% w/v solution)	6.21
Water	87.29

[0018] Azithromycin dihydrate is dispersed in the sterile unneutralised Carbopol in water containing chlorbutol BP in solution. A sterile 4% w/v sodium hydroxide solution is then added with constant mixing to a final pH of 4-6.

EXAMPLE 6

[0019] The following ingredients are formed into a mixture:

	% w/w
Azithromycin Dihydrate	3.50
Chlorbutol BP	0.50
Citric acid monohydrate**	0.117
Sodium citrate dihydrate**	0.112
Sodium citrate 1 % solution**	qs
Hydroxypropylmethylcellulose 2906 USP 4000 cps (sterile)	3.80
Water	to 100.00

** buffers

Citric acid, sodium citrate and chlorbutol BP are dissolved in 95% of the total water and the solution sterilized. Azithromycin powder is dispersed in the solution at ambient temperature using a high shear mixer. The hydroxypropylmethylcellulose, previously sterilized, is dispersed in the suspension and then allowed to hydrate over a period of about 15 minutes. The pH is adjusted to between 4-6 with a 1 % solution of sterilized sodium citrate. The gel is adjusted to final weight with water and mixed thoroughly.

EXAMPLE 7

[0020] The following suspension, gelling in situ at body temperature, is prepared:

	% w/w
Azithromycin Dihydrate	3.50
benzalkonium chloride BP	0.02
citric acid monohydrate	0.117
sodium citrate dihydrate	0.112
Pluronic® F127**	19.00
sodium citrate/citric acid solution	qs
water	to 100.00

** Pluronic® F127 is a polyoxyethylene-polyoxpropylene block copolymer of average molecular weight about 11,500

Citric acid, sodium citrate and benzalkonium chloride are dissolved in 98% of the total water. The Pluronic® F127 is dispersed in this solution and left to hydrate overnight. The preparation is then thoroughly mixed and the pH adjusted to 4-6 with sodium citrate or citric acid solution as appropriate. The solution is made to 96.5% of the total weight and sterile filtered into a sterile container. Azithromycin is dispersed aseptically in the filtered solution using a high shear mixer.

EXAMPLE 8

[0021] The following gel is prepared under strictly aseptic conditions:

	% w/w
Azithromycin Dihydrate	3.50
chlorbutol BP	0.50
ethylene maleic anhydride resin (EMA) type 91 (sterile)	0.80
dilute ammonium hydroxide solution (1.75% NH ₃)	4.40
water	90.80

The sterile EMA resin is dispersed in 50% of the total water, dilute ammonium hydroxide solution is stirred in and the mixture is heated at 95°C for 15 minutes. The resultant gel is allowed to cool to below 60°C.

[0022] The chlorbutol BP is dissolved in the remaining 50% of the water, at a temperature not exceeding 60°C, and sterile filtered into the gel which is mixed slowly.

[0023] Azithromycin is thoroughly dispersed in the gel.

Claims

1. The use of azithromycin for the manufacture of a topical medicament for treating an ocular infection.
2. The use as claimed in claim 1 wherein azithromycin is present in a composition comprising a pharmaceutically acceptable topical vehicle at a concentration of from 0.1 to 2.5 weight percent.
3. A composition for topical application directly to an eye of an animal, including a human, said composition being suitable for the treatment of an ocular infection and comprising an effective amount of azithromycin in a pharmaceutical vehicle suitable for topical application to the eye.
4. A composition as defined in claim 3 wherein azithromycin is present in a range of from 0.1 to 2.5 weight %.
5. A composition as defined in claim 4, wherein azithromycin is present in an amount of about 0.5 weight %.
6. A composition as defined in claim 5 wherein said composition is about 0.5% w/w azithromycin dihydrate suspended in an inert, non-allergenic, preservative-free vehicle comprising white petrolatum (about 55% w/w), mineral oil (about 42.5% w/w), and lanolin (about 2% w/w).



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EUROPEAN SEARCH REPORT

Application Number
EP 98 30 9274

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
X	WO 96 20010 A (ORIDIGM CORPORATION) 4 July 1996 * the whole document * * especially page 10, line 12; page 153, line 31-page 154, line 17; page 155, line 33-page 157, line 17 *	1-5	A61K31/70
D,X	US 4 474 768 A (PFIZER INC.) 2 October 1984 * the whole document * * especially column 3, line 59-column 7, line 2 *	3-5	
X	WO 95 09601 A (THE PROCTOR AND GAMBLE COMPANY) 13 April 1995 * the whole document * * especially page 7, line 17-page 8, line 4 *	3-5	
X	WO 96 19489 A (RUSSINSKY LTD.) 27 June 1996 * the whole document * * especially page 3, line 4-10 *	3-5	TECHNICAL FIELDS SEARCHED (Int.Cl.6) A61K
A	JARURATANASIRIKUL, SUTEP ET AL: "Distribution of azithromycin into brain tissue, cerebrospinal fluid, and aqueous humor of the eye" ANTIMICROB. AGENTS CHEMOTHER., 1996, 40, 825-6, XP002098516 * the whole document *	1-6	
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 30 March 1999	Examiner Mair, J
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document			

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EUROPEAN SEARCH REPORT

Application Number
EP 98 30 9274

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
A	BAILEY RL ET AL: "Randomised controlled trial of single-dose azithromycin in treatment of trachoma." LANCET, AUG 21 1993, 342 (8869) P453-6, XP002098517 ENGLAND * the whole document *	1-6	
A	THYLEFORS B.: "Azithromycin: A new opportunity for control of trachoma" WHO DRUG INFORMATION_(WHO DRUG INF.), 10/3 (132-133), XP002098518 Switzerland * the whole document *	1-6	
			TECHNICAL FIELDS SEARCHED (Int.Cl.6)
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 30 March 1999	Examiner Mair, J
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			

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**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 98 30 9274

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The members are as contained in the European Patent Office EDP file on
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30-03-1999

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9620010 A	04-07-1996	US 5872104 A	16-02-1999
		AU 4604596 A	19-07-1996
US 4474768 A	02-10-1984	AT 23343 T	15-11-1986
		AU 540056 B	01-11-1984
		AU 1692383 A	26-01-1984
		CA 1202963 A	08-04-1986
		CA 1202619 C	01-04-1986
		CA 1202620 C	01-04-1986
		CS 8305758 A	15-08-1985
		DD 215787 A	21-11-1984
		DK 332583 A,B,	20-01-1984
		EG 16882 A	30-06-1989
		EP 0101186 A	22-02-1984
		FI 832606 A	20-01-1984
		FI 864479 A,B,	04-11-1986
		GR 77556 A	24-09-1984
		IE 55365 B	29-08-1990
		JP 1193292 A	03-08-1989
		PH 18054 A	18-03-1985
		PT 77062 A,B	01-08-1983
		SU 1274626 A	30-11-1986
		JP 1036834 B	02-08-1989
		JP 1554109 C	04-04-1990
		JP 59031794 A	20-02-1984
		ZA 8305204 A	27-02-1985
WO 9509601 A	13-04-1995	AU 7957994 A	01-05-1995
		CA 2173109 A	13-04-1995
		EP 0721324 A	17-07-1996
		JP 9503504 T	08-04-1997
WO 9619489 A	27-06-1996	AU 4399396 A	10-07-1996
		IE 950955 A	26-06-1996

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